

Substitute Bill No. 7174

January Session, 2019



AN ACT CONCERNING PRESCRIPTION DRUGS.

Be it enacted by the Senate and House of Representatives in General Assembly convened:

- 1 Section 1. (NEW) (Effective October 1, 2019) (a) For the purposes of
- 2 this section:
- 3 (1) "Drug" has the same meaning as provided in section 21a-92 of 4 the general statutes;
- 5 (2) "Participating individual" means an individual resident of this 6 state who is participating in the program;
- 7 (3) "Participating pharmacist" means a pharmacist who is 8 participating in the program;
- 9 (4) "Participating pharmacy" means a pharmacy that is participating 10 in the program;
- 11 (5) "Pharmacist" has the same meaning as provided in section 38a-12 479aaa of the general statutes;
- 13 (6) "Pharmacy" has the same meaning as provided in section 38a-14 479aaa of the general statutes;
- 15 (7) "Pharmacy benefits manager" has the same meaning as provided 16 in section 38a-479aaa of the general statutes;

- 17 (8) "Program" means the Connecticut Prescription Drug Program 18 established by the Comptroller pursuant to subsection (b) of this 19 section; and
 - (9) "Program price" means the reimbursement rates and prescription drug prices established under the program.
 - (b) The Comptroller shall, within available appropriations, establish the Connecticut Prescription Drug Program. The purposes of the program shall be to: (1) Purchase outpatient prescription drugs, replenish supplies of outpatient prescription drugs or reimburse participating pharmacies and participating pharmacists for outpatient prescription drugs in order to secure the lowest possible prices and greatest possible rebates for outpatient prescription drugs prescribed to participating individuals; (2) make outpatient prescription drugs available at the lowest possible cost to participating individuals; (3) maintain a list of the most cost-effective and therapeutically effective outpatient prescription drugs available to participating individuals; (4) purchase and provide discounted outpatient prescription drugs to participating individuals; and (5) coordinate a comprehensive pharmacy benefit for participating individuals.
 - (c) (1) As part of the program, the Comptroller shall: (A) Establish eligibility criteria for individual residents of this state, as well as pharmacies and pharmacists, to participate in the program; (B) prescribe an application form for (i) individual residents of this state to become participating individuals, (ii) pharmacists to become participating pharmacists, and (iii) pharmacies to become participating pharmacies; (C) issue to participating individuals a prescription drug identification card containing the information necessary for claims processing; (D) establish a list of preferred outpatient prescription drugs for the program; (E) negotiate with pharmaceutical manufacturers and other persons to secure discounts and rebates for outpatient prescription drugs; (F) establish program prices; (G) adjudicate pharmacy claims and reimburse participating pharmacies and participating pharmacists at program prices; (H) develop a system

- for allocating and distributing the operational costs of the program, as well as any rebates, to participating individuals; and (I) charge administrative fees to participating individuals, participating pharmacists and participating pharmacies to cover the operational costs of the program, and deposit such fees in the account established under section 2 of this act.
 - (2) As part of the program, the Comptroller may: (A) Purchase outpatient prescription drugs on behalf of participating individuals; or (B) cooperate with other states or regional consortia to purchase outpatient prescription drugs on behalf of participating individuals.
 - (3) The Comptroller may enter into a contract with a pharmacy benefits manager to perform the Comptroller's duties under subdivisions (1) and (2) of this subsection, provided the Comptroller shall require the pharmacy benefits manager to charge such pharmacy benefits manager's lowest available rate to perform such duties.
- 65 (d) The Comptroller may adopt regulations, in accordance with 66 chapter 54 of the general statutes, to implement the provisions of this 67 section.
- 68 Sec. 2. (NEW) (Effective October 1, 2019) There is established an 69 account to be known as the "Connecticut Prescription Drug Program 70 account" which shall be a separate, nonlapsing account within the 71 General Fund. The account shall contain any moneys required by law 72 to be deposited in the account. Moneys in the account shall be 73 expended by the Comptroller for the purposes of the Connecticut 74 Prescription Drug Program established pursuant to section 1 of this 75 act.
- Sec. 3. (NEW) (*Effective October 1, 2019*) (a) Each pharmaceutical manufacturer doing business in this state that manufactures a brand name prescription drug and enters into an agreement with another pharmaceutical manufacturer for the purpose of delaying or preventing such other manufacturer from introducing a generic

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- 81 substitute for such drug into the marketplace shall, not later than thirty
- 82 days after entering into such agreement, send notice to the Insurance
- 83 Commissioner, in a form and manner prescribed by the commissioner,
- 84 disclosing the name of such drug.

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- (b) (1) The commissioner shall, not later than thirty days after receiving a notice pursuant to subsection (a) of this section, send notice to each health carrier, as defined in section 38a-1080 of the general statutes, and pharmacy benefits manager, as defined in section 38a-479aaa of the general statutes, doing business in this state. Such notice shall, at a minimum:
- 91 (A) Disclose the name of the brand name prescription drug that is 92 the subject of the notice the commissioner received pursuant to 93 subsection (a) of this section; and
 - (B) Instruct such health carrier, if such health carrier includes such drug on such health carrier's drug formulary or list of covered drugs, or pharmacy benefits manager, if such pharmacy benefits manager administers a prescription drug benefit that includes such drug, to immediately reduce the cost of such drug to covered individuals by an amount that is equal to fifty per cent of the manufacturer's wholesale list price for such drug.
- 101 (2) For the purposes of this subsection, "manufacturer's wholesale 102 list price" has the same meaning as provided in section 21a-126 of the 103 general statutes.
- 104 (c) The provisions of this section shall apply to the maximum extent permitted by applicable law.
- 106 (d) The commissioner may adopt regulations, in accordance with 107 chapter 54 of the general statutes, to implement the provisions of this 108 section.
- Sec. 4. Subdivision (3) of subsection (m) of section 5-259 of the general statutes is repealed and the following is substituted in lieu

th	nereof (Effective October 1, 2019):
	(3) (A) [(i)] For the purposes of this subdivision:
	(i) "Nonstate public employer" means (I) a municipality or other
po	olitical subdivision of the state, including a board of education, quasi-
_	ublic agency or public library, as defined in section 11-24a, or (II) the
_	eachers' Retirement Board; and
	(ii) "Qualified private employer" means a self-insured private
<u>er</u>	mployer doing business in this state.
	(B) The Comptroller shall offer nonstate public employers and
<u>q</u> 1	ualified private employers the option to purchase prescription drugs
fc	or their employees, employees' dependents and retirees under the
pı	urchasing authority of the state pursuant to section 1 of public act 09-
20	06, subject to the provisions of subparagraph [(E)] (F) of this
SU	abdivision.
	[(ii) For purposes of this subdivision, "nonstate public employer"
m	eans (I) a municipality or other political subdivision of the state,
in	cluding a board of education, quasi-public agency or public library,
as	s defined in section 11-24a, or (II) the Teachers' Retirement Board.]
	[(B)] (C) The Comptroller shall establish procedures to determine (i)
th	ne eligibility requirements for, (ii) the enrollment procedures for, (iii)
th	ne duration of, (iv) requirements regarding payment for, and (v) the
pı	rocedures for withdrawal from and termination of, the purchasing of
pı	rescription drugs for nonstate public employers and qualified private
<u>er</u>	<u>mployers</u> under subparagraph $[(A)]$ (B) of this subdivision.
	[(C)] (D) The Comptroller may offer to nonstate public employers
ar	nd qualified private employers that choose to purchase prescription
dı	rugs pursuant to subparagraph [(A)] (B) of this subdivision the
01	ption to purchase stop loss coverage from an insurer at a rate

negotiated by the Comptroller.

- [(D)] (E) Two or more nonstate public employers or qualified private employers may join together for the purpose of purchasing prescription drugs for their employees, employees' dependents and retirees. Such arrangement shall not constitute a multiple employer welfare arrangement, as defined in Section 3 of the Employee Retirement Income Security Act of 1974, as amended from time to time.
- 147 [(E)] (F) (i) The Comptroller shall offer nonstate public employers 148 and qualified private employers the option to purchase prescription 149 drugs through the plan set forth in the State Employees' Bargaining 150 Agent Coalition's collective bargaining agreement with the state only if 151 the Health Care Cost Containment Committee, established in 152 accordance with the ratified agreement between the state and said 153 coalition pursuant to subsection (f) of section 5-278, has indicated in 154 writing to the Comptroller that allowing such nonstate public 155 employers and qualified private employers such option is consistent 156 with said coalition's collective bargaining agreement.
- (ii) Such writing shall not be required if the Comptroller establishes a separate prescription drugs purchasing plan <u>or plans</u> for nonstate public employers <u>and qualified private employers</u>.
- (iii) Nonstate public employers <u>and qualified private employers</u> that purchase prescription drugs pursuant to this subdivision shall pay the full cost of their own claims and prescription drugs.
- Sec. 5. Section 38a-477cc of the general statutes is repealed and the following is substituted in lieu thereof (*Effective October 1, 2019*):
- (a) [On and after January 1, 2018, no] <u>No</u> contract for pharmacy services entered into in the state between a health carrier, as defined in section 38a-591a, or pharmacy benefits manager, as defined in section 38a-479aaa, and a pharmacy or pharmacist shall:
- 169 (1) On and after January 1, 2018, contain a provision prohibiting or penalizing, including through increased utilization review, reduced

- payments or other financial disincentives, a pharmacist's disclosure to an individual purchasing prescription medication of information regarding: [(1) the] (A) The cost of the prescription medication to the individual; [,] or [(2) the] (B) The availability of any therapeutically equivalent alternative medications or alternative methods of purchasing the prescription medication, including, but not limited to, paying a cash price, that are less expensive than the cost of the prescription medication to the individual; [.] and
- (2) On and after January 1, 2020, contain a provision permitting the health carrier or pharmacy benefits manager to recoup, directly or indirectly, from a pharmacy or pharmacist any portion of a claim that such health carrier or pharmacy benefits manager has paid to the pharmacy or pharmacist, unless such recoupment is permitted under section 38a-479iii or required by applicable law.
 - (b) (1) On and after January 1, 2018, no health carrier or pharmacy benefits manager shall require an individual to make a payment at the point of sale for a covered prescription medication in an amount greater than the lesser of:
 - [(1) the] (A) The applicable copayment for such prescription medication; [, (2) the] (B) The allowable claim amount for the prescription medication; [,] or [(3) the] (C) The amount an individual would pay for the prescription medication if the individual purchased the prescription medication without using a health benefit plan, as defined in section 38a-591a, or any other source of prescription medication benefits or discounts.
 - (2) For the purposes of this subsection, "allowable claim amount" means the amount the health carrier or pharmacy benefits manager has agreed to pay the pharmacy for the prescription medication.
 - (c) Any provision of a contract that violates the provisions of this section shall be void and unenforceable. Any general business practice that violates the provisions of this section shall constitute an unfair

- 202 trade practice pursuant to chapter 735a. The invalidity or
- 203 unenforceability of any contract provision under this subsection shall
- 204 not affect any other provision of the contract.
- 205 (d) The Insurance Commissioner may: [, (1) pursuant to the 206 provisions of chapter 697, enforce]
- 207 (1) Enforce the provisions of this section [,] pursuant to chapter 697; 208 and
- 209 (2) [upon] <u>Upon</u> request, audit a contract for pharmacy services for compliance with the provisions of this section.
- Sec. 6. (*Effective from passage*) (a) There is established a task force to
- 212 study drug reimportation. Such study shall include, but need not be
- 213 limited to, an examination of the feasibility of implementing a drug
- 214 reimportation program for the purpose of lowering the cost of
- 215 prescription drugs and health insurance in this state.
- 216 (b) The task force shall consist of the following members:
- 217 (1) Two appointed by the speaker of the House of Representatives;
- 218 (2) Two appointed by the president pro tempore of the Senate;
- 219 (3) One appointed by the majority leader of the House of
- 220 Representatives;
- 221 (4) One appointed by the majority leader of the Senate;
- 222 (5) One appointed by the minority leader of the House of
- 223 Representatives;
- 224 (6) One appointed by the minority leader of the Senate;
- 225 (7) The Attorney General, or the Attorney General's designee;
- 226 (8) The Comptroller, or the Comptroller's designee;

227 (9) The Insurance Commissioner, or the commissioner's designee; 228 (10) The Commissioner of Public Health, or the commissioner's 229 designee; 230 (11) The Commissioner of Social Services, or the commissioner's 231 designee; 232 (12) The executive director of the Office of Health Strategy, or the 233 executive director's designee; 234 (13) The Healthcare Advocate, or the Healthcare Advocate's 235 designee; and 236 (14) Two persons appointed by the Governor. 237 (c) Any member of the task force appointed under subdivision (1), 238 (2), (3), (4), (5) or (6) of subsection (b) of this section may be a member 239 of the General Assembly. 240 (d) All appointments to the task force shall be made not later than 241 thirty days after the effective date of this section. Any vacancy shall be 242 filled by the appointing authority. 243 (e) The speaker of the House of Representatives and the president 244 pro tempore of the Senate shall select the chairpersons of the task force 245 from among the members of the task force. Such chairpersons shall 246 schedule the first meeting of the task force, which shall be held not 247 later than sixty days after the effective date of this section. 248 (f) The administrative staff of the joint standing committee of the 249 General Assembly having cognizance of matters relating to insurance 250 shall serve as administrative staff of the task force. 251 (g) Not later than January 1, 2020, the task force shall submit a 252 report on its findings and recommendations to the joint standing 253 committee of the General Assembly having cognizance of matters

relating to insurance, in accordance with the provisions of section 11-

4a of the general statutes. The task force shall terminate on the date that it submits such report or January 1, 2020, whichever is later.

This act shall take effect as follows and shall amend the following				
sections:				
Section 1	October 1, 2019	New section		
Sec. 2	October 1, 2019	New section		
Sec. 3	October 1, 2019	New section		
Sec. 4	October 1, 2019	5-259(m)(3)		
Sec. 5	October 1, 2019	38a-477cc		
Sec. 6	from passage	New section		

Statement of Legislative Commissioners:

In Section 3(b)(2), "subdivision" was changed to "subsection" for accuracy.

INS Joint Favorable Subst. -LCO